Certificate of CE-Registration

This is to certify that, in accordance with the *In Vitro* Diagnostic Medical Device Directive 98/79/EC and the Medical Device Directive 93/42/EEC, Medical Device Safety Service GmbH (MDSS) agrees to perform all duties and responsibilities as the Authorized Representative for:

Biomiga Inc.
10637 Roselle St Suite C
San Diego, CA 92121
USA

as stipulated and demanded by the aforementioned Directive. The European Databank on Medical Devices (EUDAMED) is established as of May 1, 2011. The German Competent Authority is notified of the manufacturer's *in vitro* diagnostic medical devices and medical devices and has allocated registration numbers shown in:

Annex A: February 22, 2021

The Manufacturer has provided MDSS with the appropriate Declaration(s) of Conformity confirming that the *in vitro* diagnostic medical devices and medical devices fulfill the applicable requirements of Directives 98/79/EC and 93/42/EEC. In compliance with German law, a safety officer has been appointed for Germany.

2021-02-22

Dr. Philipp Hohenbrink
Senior Consultant
MDSS GmbH

Annex A: February 22, 2021
Manufacturer: Biomiga Inc.

Model, Reference, Catalog or UDI Number	Device Names (notified)	Nomenclature (notified)	Code (notified)	Description (notified)	Class (notified)	EC Certificate No. & Expiry (notified)	German Registration Number
VR6568	Viral RNA Extraction from Respiratory Specimens	GMDN	52521	Nucleic acid extraction/isolation kit IVD	"Other"	N.A.	DE/CA09/0170/B29/IVD/001
MR6532	MgPure Viral RNA Purification Kit	GMDN	52521	Nucleic acid extraction/isolation kit IVD	"Other"	N.A.	DE/CA09/0170/B29/IVD/001